

**Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings of claims in the application:

**Listing of Claims:**

1. (Previously presented) A method for isolating an analyte contained in a gas phase comprising the steps of:
  - a) providing device comprising a gas- and liquid-permeable carrier matrix containing a first binding partner of an analyte in elutable form, and
  - b) contacting a gas phase suspected of containing said analyte with said gas- and liquid-permeable carrier matrix containing a first binding partner of an analyte in elutable form such that said analyte binds to said first binding partner, and
  - c) eluting any complex consisting of said analyte and said first binding partner and any uncomplexed first binding partner from said device, and
  - d) isolating said analyte.
2. (Previously presented) The method according to claim 1, wherein after step c) said analyte is released from said complex consisting of said analyte and said first binding partner.
3. (Currently Amended) A method for immunologically detecting an analyte contained in a gas phase, comprising the steps of:
  - a) providing a device comprising: a gas- and liquid-permeable carrier matrix containing a first binding partner of an analyte in elutable form,

- b) contacting a sample gas suspected of containing said analyte said gas- and liquid-permeable carrier matrix containing a first binding partner of an analyte in elutable form such that said analyte binds to said first binding partner,
- c) eluting any complex consisting of said analyte and said first binding partner and any uncomplexed first binding partner from said first carrier matrix, and
- d) determining any eluted complex ~~of~~ or uncomplexed first binding partner as a measure of the amount of analyte present.

- 4. (Canceled)
- 5. (Original) The method according to claim 3, further comprising labeling the first binding partner, and determining any labeled eluted complex or labeled uncomplexed first binding as a measure of the amount of analyte present.
- 6. (Original) The method according to claim 5, wherein said first binding partner is enzymatically labeled.
- 7. (Original) Method according to claim 3, further comprising,
  - a) binding any eluted first binding partner to a labeled second binding partner specific for said first binding partner, and
  - b) determining any bound label as a measure of the amount of analyte present, wherein said first binding partner is unlabeled.

8. (Original) The method according to claim 7, wherein said first binding partner is bound to said labeled second binding partner in a second carrier matrix and said determination of said label is in a third carrier matrix.
9. (Original) The method according to claim 1, wherein said first carrier matrix has a liquid content of 10 to 90%.
10. (Original) The method according to claim 3, wherein said first carrier matrix has liquid content of 10 to 90%.
11. (Original) The method according to claim 9, wherein said analyte is first nonspecifically absorbed to an essentially dry matrix and then a liquid is added allowing an immune reaction between said analyte and said first binding partner to occur.
12. (Original) The method according to claim 10, wherein said analyte is first nonspecifically absorbed to an essentially dry matrix and then a liquid is added allowing an immune reaction between said analyte and said first binding partner to occur.
13. (Original) The method according to claim 9, wherein said liquid is an aqueous solution containing 0 to 30% polar organic solvent and 0 to 1% detergent.

14. (Original) The method according to claim 10, wherein said liquid is an aqueous solution containing 0 to 30% polar organic solvent and 0 to 1% detergent.
15. (Original) The method according to claim 3, wherein said first binding partner is an antibody or Fab' fragment thereof.
16. (Original) The method according to claim 3, wherein said first binding partner is a receptor which binds said analyte.
17. (Original) The method according to claim 3, wherein said first carrier matrix is a hydrophilic or hygroscopic material.
18. (Original) The method according to claim 17, wherein said hydrophilic or hygroscopic material is the form of particles or fibers.

Claims 19-22 (Canceled).

23. (Currently Amended) A method for detecting the presence of an analyte ~~of a substance~~ in a gas permeable, enclosed package, comprising the steps of:
  - a) drawing a sample of a gas surrounding a gas permeable, enclosed package suspected of containing an analyte ~~of a substance~~ of interest,

- b) contacting a first carrier matrix with said sample of gas, wherein a first binding partner, specific for ~~an~~ said analyte ~~of said substance~~, is contained in said first carrier matrix but not bound to said first carrier matrix,
  - c) binding any analyte ~~contained in said substance~~ present in said sample of gas to said first binding partner,
  - d) eluting any complex consisting of said analyte and said first binding partner and any uncomplexed first binding partner from said first carrier matrix, and
  - e) determining any eluted complex or uncomplexed first binding partner as a measure of the presence of said analyte ~~of said substance~~.
24. (Currently amended) The method according to claim 23, wherein said ~~substance~~ analyte is selected from the group consisting of nitroglycol, nitroglycerin, nitropenta, hexogen, octogen, tetranitromethane, trinitrotoluene, trinitrobenzene, trinitroanisol, triaminotrinitrotolulene, hexanitrostilben, polycyclic aromatic hydrocarbons, polychlorated biphenyls, herbicides and pesticides.
25. (Currently amended) The method according to claim 23, wherein said ~~substance~~ analyte is an illegal drug of abuse.
26. (Currently amended) The method according to claim 25, wherein said ~~substance~~ analyte is selected from the group consisting of cocaine, heroin, cannabinol, cannabidiol and tetrahydrocannabinol.

27. (Original) The method according to claim 23, wherein said enclosed package is luggage.
28. (Previously presented) The method according to claim 4, wherein said first carrier matrix has a gas permeability between 10 ml/min to 100 l/min.
29. (Previously presented) The method according to claim 28, wherein said first carrier matrix has a gas permeability between 100 ml/min to 20 l/min.
30. (Previously presented) The method according to claim 29, wherein said first carrier matrix has a gas permeability between 500 ml/min to 10 l/min.